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NEWS	1		Web Page for STN Seminar Schedule - N. America
NEWS	2	APR 04	STN AnaVist, Version 1, to be discontinued
NEWS	3	APR 15	WPIDS, WPINDEX, and WPIX enhanced with new predefined hit display formats
NEWS	4	APR 28	EMBASE Controlled Term thesaurus enhanced
NEWS	5	APR 28	IMSRESEARCH reloaded with enhancements
NEWS	6	MAY 30	INPAFAMDB now available on STN for patent family searching
NEWS	7	MAY 30	DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
NEWS	8	JUN 06	EPFULL enhanced with 260,000 English abstracts
NEWS	9	JUN 06	KOREAPAT updated with 41,000 documents
NEWS	10	JUN 13	USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
NEWS	11	JUN 19	CAS REGISTRY includes selected substances from web-based collections
NEWS	12	JUN 25	CA/CAPplus and USPAT databases updated with IPC reclassification data
NEWS	13	JUN 30	AEROSPACE enhanced with more than 1 million U.S. patent records
NEWS	14	JUN 30	EMBASE, EMBAL, and LEMBASE updated with additional options to display authors and affiliated organizations
NEWS	15	JUN 30	STN on the Web enhanced with new STN AnaVist Assistant and BLAST plug-in
NEWS	16	JUN 30	STN AnaVist enhanced with database content from EPFULL
NEWS	17	JUL 28	CA/CAPplus patent coverage enhanced
NEWS	18	JUL 28	EPFULL enhanced with additional legal status information from the epline Register
NEWS	19	JUL 28	IFICDB, IFIPAT, and IFIUIDB reloaded with enhancements
NEWS	20	JUL 28	STN Viewer performance improved
NEWS	21	AUG 01	INPADOCDB and INPAFAMDB coverage enhanced
NEWS	22	AUG 13	CA/CAPplus enhanced with printed Chemical Abstracts page images from 1967-1998
NEWS	23	AUG 15	CAOLD to be discontinued on December 31, 2008
NEWS	24	AUG 15	CAPplus currency for Korean patents enhanced
NEWS	25	AUG 25	CA/CAPplus, CASREACT, and IFI and USPAT databases enhanced for more flexible patent number searching
NEWS	26	AUG 27	CAS definition of basic patents expanded to ensure comprehensive access to substance and sequence information

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,  
AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

NEWS HOURS STN Operating Hours Plus Help Desk Availability

NEWS LOGIN      Welcome Banner and News Items  
NEWS IPC8        For general information regarding STN implementation of IPC 8

Enter NEWS followed by the item number or name to see news on that specific topic.

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\* \* \* \* \* STN Columbus \* \* \* \* \*

FILE 'HOME' ENTERED AT 10:29:53 ON 17 SEP 2008

=> file reg		
COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	0.21	0.21

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STRUCTURE FILE UPDATES: 16 SEP 2008 HIGHEST RN 1049663-83-3  
DICTIONARY FILE UPDATES: 16 SEP 2008 HIGHEST RN 1049663-83-3

New CAS Information Use Policies, enter HELP USAGETERMS for details.

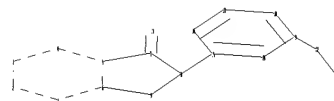
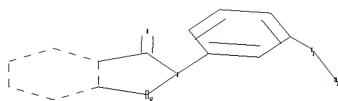
TSCA INFORMATION NOW CURRENT THROUGH July 5, 2008.

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

=>  
Uploading C:\Program Files\Stnexp\Queries\10551904.str



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chain nodes :
10 11 22 23
ring nodes :
1 2 3 4 5 6 7 8 9 15 16 17 18 19 20
chain bonds :
7-10 8-15 19-22 22-23
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-9 7-8 8-9 15-16 15-20 16-17 17-18 18-19
19-20
exact/norm bonds :
1-2 1-6 2-3 3-4 4-5 5-6 7-8 8-9 8-15 19-22 22-23
exact bonds :
5-7 6-9 7-10
normalized bonds :
15-16 15-20 16-17 17-18 18-19 19-20
isolated ring systems :
containing 1 : 15 :

```

G1:C,O,S,N

```

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:CLASS 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom 20:Atom 22:CLASS 23:CLASS

```

L1           STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1           STR

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 10:30:35 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED -           16 TO ITERATE

100.0% PROCESSED           16 ITERATIONS

7 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:   ONLINE   \*\*COMPLETE\*\*

BATCH   \*\*COMPLETE\*\*

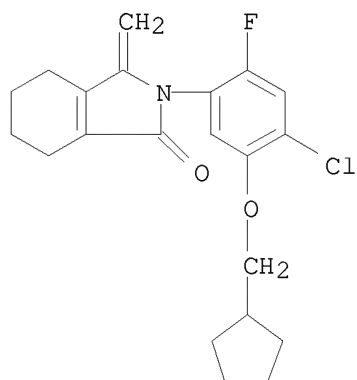
PROJECTED ITERATIONS:           80 TO           560

PROJECTED ANSWERS:               7 TO           298

L2               7 SEA SSS SAM L1

=> d scan

L2 7 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN  
IN 1H-Isoindol-1-one, 2-[4-chloro-5-(cyclopentylmethoxy)-2-fluorophenyl]-  
2,3,4,5,6,7-hexahydro-3-methylene-  
MF C21 H23 Cl F N O2



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> s l1 full  
FULL SEARCH INITIATED 10:30:45 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 173 TO ITERATE

100.0% PROCESSED 173 ITERATIONS 45 ANSWERS  
SEARCH TIME: 00.00.01

L3 45 SEA SSS FUL L1

=> file caplus  
COST IN U.S. DOLLARS SINCE FILE TOTAL  
ENTRY SESSION  
FULL ESTIMATED COST 178.36 178.57

FILE 'CAPLUS' ENTERED AT 10:30:49 ON 17 SEP 2008  
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FILE COVERS 1907 - 17 Sep 2008 VOL 149 ISS 12  
FILE LAST UPDATED: 16 Sep 2008 (20080916/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

=> s l3 full  
L4 10 L3

=> d ibib abs hitstr tot

L4 ANSWER 1 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:486256 CAPLUS

DOCUMENT NUMBER: 139:261118

TITLE: New reaction of ethenetetracarbonitrile with  
N-arylisindolines

AUTHOR(S): Dopp, Dietrich; Hassan, Alaa A.; Mourad, Aboul-Fetouh  
E.; Nour El-Din, Ahmed M.; Angermund, Klaus; Kruger,  
Carl; Lehmann, Christian W.; Rust, Jorg

CORPORATE SOURCE: Institut fur Chemie, Universitat Duisburg-Essen,  
Duisburg, D-47048, Germany

SOURCE: Tetrahedron (2003), 59(27), 5073-5081  
CODEN: TETRAB; ISSN: 0040-4020

PUBLISHER: Elsevier Science B.V.

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 139:261118

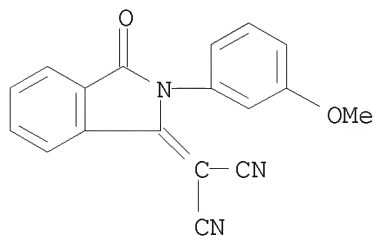
AB N-arylisindolines react with ethenetetracarbonitrile in aerated benzene by formation of [3-(2-aryl-3-dicyanomethylene-2,3-dihydro-1H-indol-1-ylidene)-2-aryl-2,3-dihydro-1H-indol-1-ylidene]propanedinitriles (20-36%), N-aryl-3-dicyanomethylene-indol-2-ones (15-21%) and N-arylphthalimides (4-9%) as well as 1,1,2,2-tetracyanoethane (35-55%). The structure of [3-[3-(dicyanomethylene)-2,3-dihydro-2-(3-methylphenyl)-1H-indol-1-ylidene]-2,3-dihydro-2-(3-methylphenyl)-1H-indol-1-ylidene]propanedinitrile has been unambiguously confirmed by a single crystal X-ray structure anal. A rationale for the formation of products 8-11 is presented. For example, the reaction of ethenetetracarbonitrile with 2,3-dihydro-2-(2-methoxyphenyl)-1H-indole gave [3-[3-(dicyanomethylene)-2,3-dihydro-2-(2-methoxyphenyl)-1H-indol-1-ylidene]-2,3-dihydro-2-(2-methoxyphenyl)-1H-indol-1-ylidene]propanedinitrile, [2,3-dihydro-2-(2-methoxyphenyl)-3-oxo-1H-indol-1-ylidene]propanedinitrile, 2-(2-methoxyphenyl)-1H-indole-1,3(2H)-dione, and 1,1,2,2-ethanetetracarbonitrile.

IT 170300-33-1P, [2,3-Dihydro-2-(3-methoxyphenyl)-3-oxo-1H-indol-1-ylidene]propanedinitrile 600724-02-5P

RL: SPN (Synthetic preparation); PREP (Preparation)  
(reaction of ethenetetracarbonitrile with N-arylisindolines)

RN 170300-33-1 CAPLUS

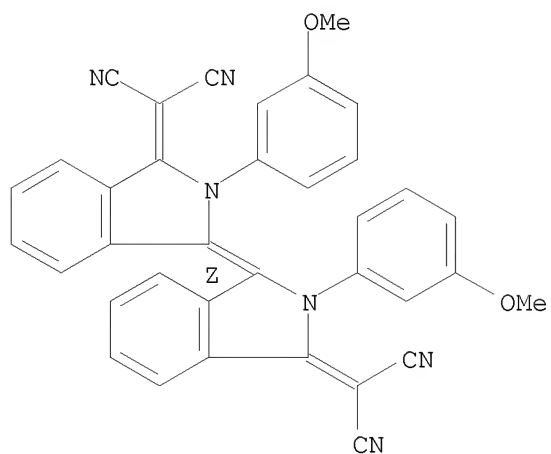
CN Propanedinitrile, 2-[2,3-dihydro-2-(3-methoxyphenyl)-3-oxo-1H-indol-1-ylidene]- (CA INDEX NAME)



RN 600724-02-5 CAPLUS

CN Propanedinitrile, [(3Z)-3-[3-(dicyanomethylene)-2,3-dihydro-2-(3-methoxyphenyl)-1H-indol-1-ylidene]-2,3-dihydro-2-(3-methoxyphenyl)-1H-indol-1-ylidene]- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



REFERENCE COUNT:

39

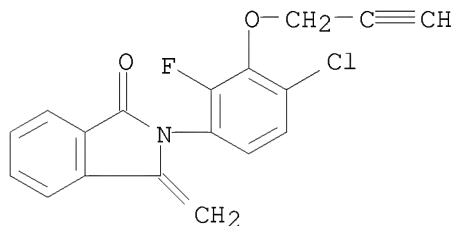
THERE ARE 39 CITED REFERENCES AVAILABLE FOR THIS  
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT



L4 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

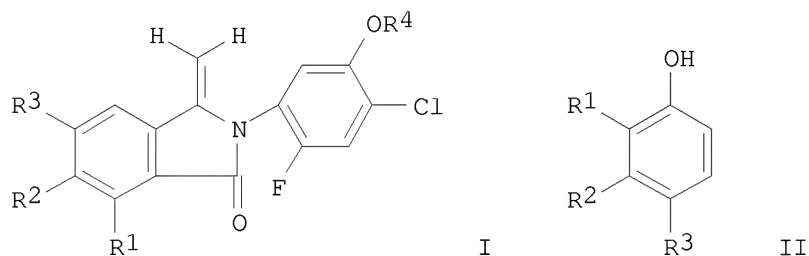
ACCESSION NUMBER: 2000:744988 CAPLUS  
DOCUMENT NUMBER: 133:266725  
TITLE: Preparation of herbicidal 2,3-dihydro-3-methylene-2-(substituted)-1H-isoindol-1-one derivatives  
INVENTOR(S): Yu, Eung-Kul; Kim, Jae-Nyung  
PATENT ASSIGNEE(S): Korea Research Institute of Chemical Technology, S. Korea  
SOURCE: Repub. Korea, No pp. given  
CODEN: KRXXFC  
DOCUMENT TYPE: Patent  
LANGUAGE: Korean  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	KR 128544	B1	19980404	KR 1994-16984	19940714
PRIORITY APPLN. INFO.:				KR 1994-16984	19940714
AB	The title derivs. [e.g., 2,3-dihydro-3-methylene-2- (3-propargyloxy-4-chloro-2-fluorophenyl)-1H-isoindol-1-one], useful as herbicides, were prepared				
IT	298185-08-7P RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of herbicidal 2,3-dihydro-3-methylene-2-(substituted)-1H-isoindol-1-one derivs.)				
RN	298185-08-7 CAPLUS				
CN	1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-3-(2-propyn-1-yloxy)phenyl]-2,3-dihydro-3-methylene- (CA INDEX NAME)				

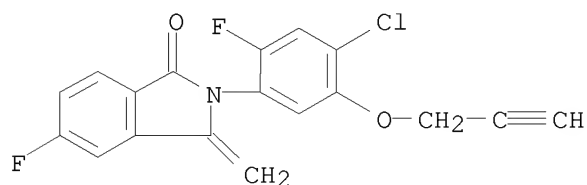


L4 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

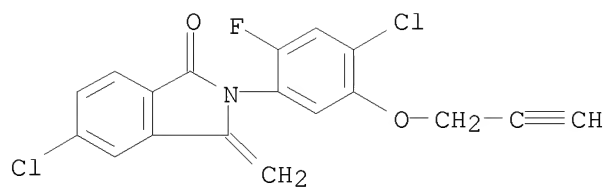
ACCESSION NUMBER: 1996:88798 CAPLUS  
DOCUMENT NUMBER: 124:232184  
ORIGINAL REFERENCE NO.: 124:43007a, 43010a  
TITLE: A facile synthetic method of herbicidal  
2,3-dihydro-3-methylene-2-substituted-phenyl-1H-  
isoindol-1-one derivatives  
AUTHOR(S): Kim, Jae Nyoung; Ryu, Eung K.  
CORPORATE SOURCE: Dep. Chem., Chonnam Natl. Univ., Kwangju, 500-757, S.  
Korea  
SOURCE: Synthetic Communications (1996), 26(1), 67-74  
CODEN: SYNCAV; ISSN: 0039-7911  
PUBLISHER: Dekker  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
OTHER SOURCE(S): CASREACT 124:232184  
GI



AB Herbicidal 2,3-dihydro-3-methylene-2-substituted phenyl-1H-isoindol-1-one  
derivs. I (R1 = H, Cl, R2 = H, OMe, Me, R3 = F, Cl, H, Me, R4 =  
CH2C.tplbond.CH, Et, CHMeCN) have been synthesized. They were prepared from  
the easily available phenol derivs. II in 5 steps in moderate yields.  
IT 174813-47-9P 174813-48-0P 174813-49-1P  
174813-50-4P 174813-51-5P 174813-52-6P  
174813-53-7P 174813-54-8P  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of methylenephénylisoindolones)  
RN 174813-47-9 CAPLUS  
CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-propyn-1-yloxy)phenyl]-5-  
fluoro-2,3-dihydro-3-methylene- (CA INDEX NAME)

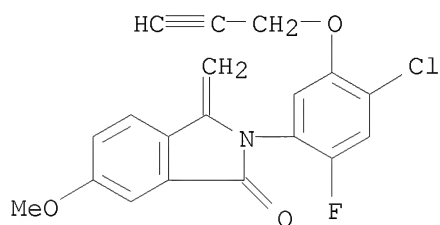


RN 174813-48-0 CAPLUS  
CN 1H-Isoindol-1-one, 5-chloro-2-[4-chloro-2-fluoro-5-(2-propyn-1-  
yloxy)phenyl]-2,3-dihydro-3-methylene- (CA INDEX NAME)



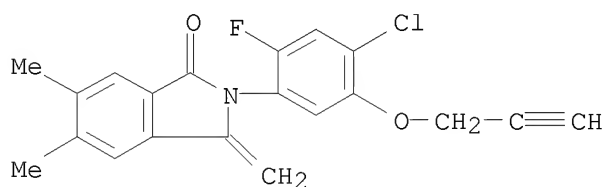
RN 174813-49-1 CAPLUS

CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-propyn-1-yloxy)phenyl]-2,3-dihydro-6-methoxy-3-methylene- (CA INDEX NAME)



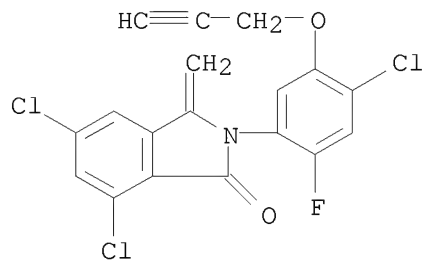
RN 174813-50-4 CAPLUS

CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-propyn-1-yloxy)phenyl]-2,3-dihydro-5,6-dimethyl-3-methylene- (CA INDEX NAME)



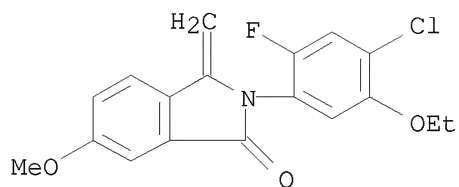
RN 174813-51-5 CAPLUS

CN 1H-Isoindol-1-one, 5,7-dichloro-2-[4-chloro-2-fluoro-5-(2-propyn-1-yloxy)phenyl]-2,3-dihydro-3-methylene- (CA INDEX NAME)



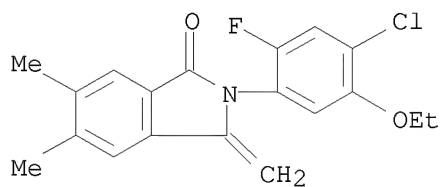
RN 174813-52-6 CAPLUS

CN 1H-Isoindol-1-one, 2-(4-chloro-5-ethoxy-2-fluorophenyl)-2,3-dihydro-6-methoxy-3-methylene- (CA INDEX NAME)



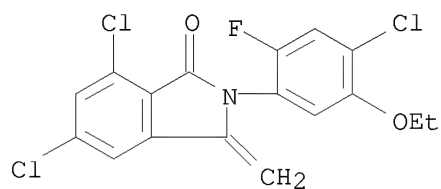
RN 174813-53-7 CAPLUS

CN 1H-Isoindol-1-one, 2-(4-chloro-5-ethoxy-2-fluorophenyl)-2,3-dihydro-5,6-dimethyl-3-methylene- (CA INDEX NAME)



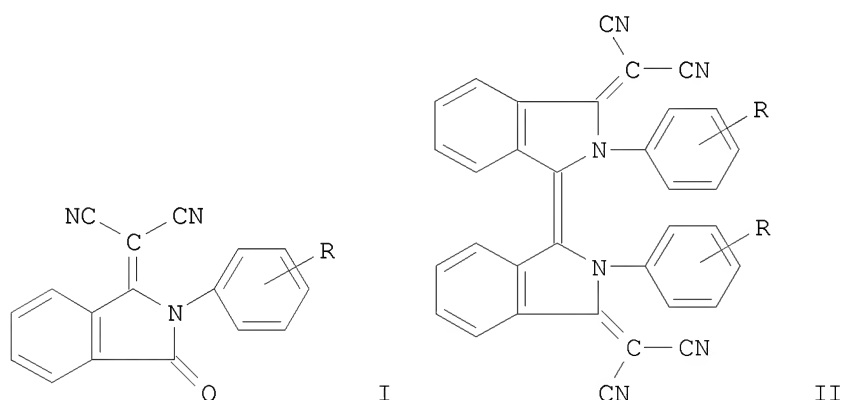
RN 174813-54-8 CAPLUS

CN 1H-Isoindol-1-one, 5,7-dichloro-2-(4-chloro-5-ethoxy-2-fluorophenyl)-2,3-dihydro-3-methylene- (CA INDEX NAME)



L4 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1995:689220 CAPLUS  
DOCUMENT NUMBER: 123:338851  
ORIGINAL REFERENCE NO.: 123:60811a,60814a  
TITLE: Chemical and electrochemical reduction of the products  
from the reactions of isoindolines and  
tetracyanoethylene  
AUTHOR(S): Carloni, Patricia; Greci, Lucedio; Stipa, Pierluigi;  
Doepp, Dietrich; Hassan, Alaa El-Din Abdel Hafeez;  
Alberti, Angelo  
CORPORATE SOURCE: Dip. Sci. Mater. Terra, Univ. Ancona, Ancona, I-60131,  
Italy  
SOURCE: Tetrahedron (1995), 51(27), 7451-8  
CODEN: TETRAB; ISSN: 0040-4020  
PUBLISHER: Pergamon  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
GI



AB The reduction potentials of compds. resulting from the title reaction and containing one (I; R=p-, m-MeO) or two (II; R=H,p-,m-MeO, p-Me) dicyanovinylidene moieties have been determined by means of cyclic voltammetry. Electrochem. reduction of the same compds. within the cavity of an EPR spectrometer led to the observation of radical species tentatively identified as the corresponding radical anions. EPR expts. have also provided indication that the first formed species in the butoxide reduction of II undergoes cleavage of the bond between the two heterocyclic systems evolving to radical anions identical to those obtained by similar reduction of compds. I. Some models compds. have also been investigated for comparison purposes.

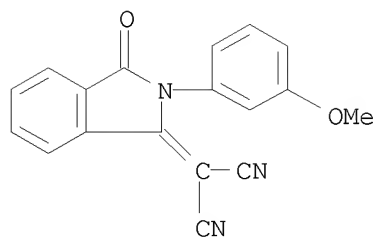
IT 170300-36-4

RL: FMU (Formation, unclassified); PRP (Properties); FORM (Formation, nonpreparative)

(ESR parameters for radical anions observed upon chemical or electrochem. reduction of isoindoline-tetracyanoethylene adducts)

RN 170300-36-4 CAPLUS

CN Propanedinitrile, [2,3-dihydro-2-(3-methoxyphenyl)-3-oxo-1H-isoindol-1-ylidene]-, radical ion(1-) (9CI) (CA INDEX NAME)

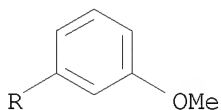
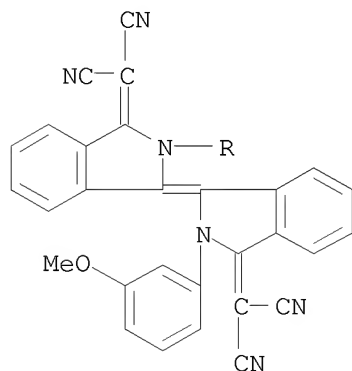


IT 170300-28-4 170300-33-1

RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)  
(ESR parameters for radical anions observed upon chemical or electrochem. reduction of isoindoline-tetracyanoethylene adducts)

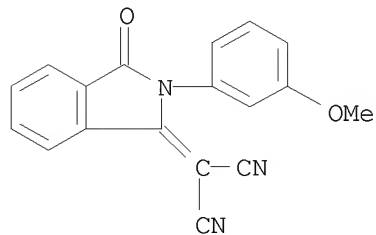
RN 170300-28-4 CAPLUS

CN Propanedinitrile, [3-[3-(dicyanomethylene)-2,3-dihydro-2-(3-methoxyphenyl)-1H-isoindol-1-ylidene]-2,3-dihydro-2-(3-methoxyphenyl)-1H-isoindol-1-ylidene]- (9CI) (CA INDEX NAME)



RN 170300-33-1 CAPLUS

CN Propanedinitrile, 2-[2,3-dihydro-2-(3-methoxyphenyl)-3-oxo-1H-isoindol-1-ylidene]- (CA INDEX NAME)

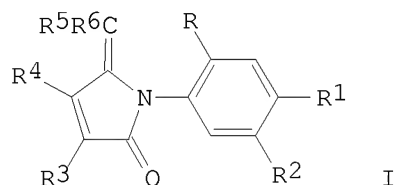


L4 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1991:207016 CAPLUS  
DOCUMENT NUMBER: 114:207016  
ORIGINAL REFERENCE NO.: 114:34911a,34914a  
TITLE: Herbicidal N-arylpyrrolinones  
INVENTOR(S): Elbe, Hans Ludwig; Marhold, Albrecht; Luerksen, Klaus;  
Santel, Hans Joachim; Schmidt, Robert R.; Krauskopf,  
Birgit  
PATENT ASSIGNEE(S): Bayer A.-G., Germany  
SOURCE: Eur. Pat. Appl., 64 pp.  
CODEN: EPXXDW  
DOCUMENT TYPE: Patent  
LANGUAGE: German  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 403891	A1	19901227	EP 1990-110853	19900608
R: BE, CH, DE, FR, GB, IT, LI, NL				
DE 3920271	A1	19910103	DE 1989-3920271	19890621
US 5045108	A	19910903	US 1990-534383	19900607
JP 03031256	A	19910212	JP 1990-158848	19900619
PRIORITY APPLN. INFO.:			DE 1989-3920271	A 19890621
OTHER SOURCE(S):	CASREACT 114:207016; MARPAT 114:207016			

GI



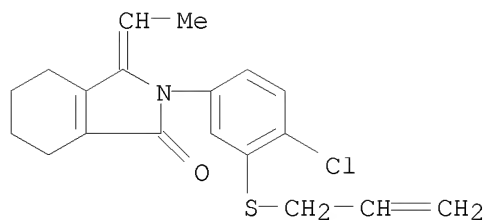
AB Title compds. I [R = H, halogen; R<sup>1</sup> = cyano, NO<sub>2</sub>, F, Cl, Br, iodo, alkyl, haloalkyl, alkoxy, haloalkoxy, alkylthio, haloalkylthio; R<sup>2</sup> = halogen, OH, SH, (un)substituted alkoxy, alkenyloxy, alkynyloxy, cycloalkoxy, alkylthio, alkenylthio, alkynylthio, cycloalkylthio; R<sup>3</sup>, R<sup>4</sup> = H, halogen, alkyl; R<sup>5</sup>, R<sup>6</sup> = H, alkyl; R<sup>3</sup>R<sup>4</sup>, R<sup>5</sup>R<sup>6</sup> = alkanediyl] were prepared Thus, MeCO<sub>2</sub>Me<sub>2</sub>CH:CHCO<sub>2</sub>H was treated with 4,2,5-NC(F<sub>2</sub>)C<sub>6</sub>H<sub>2</sub>NH<sub>2</sub> to give I (R = R<sup>2</sup> = F, R<sup>1</sup> = cyano, R<sup>3</sup> = R<sup>4</sup> = Me, R<sup>5</sup> = R<sup>6</sup> = H). I have herbicidal activity both pre- and post-emergence.

IT 133715-52-3P 133715-56-7P 133715-58-9P  
133715-60-3P 133715-62-5P 133715-63-6P  
133715-64-7P 133715-66-9P 133715-69-2P  
133715-71-6P 133715-72-7P 133715-73-8P  
133715-74-9P

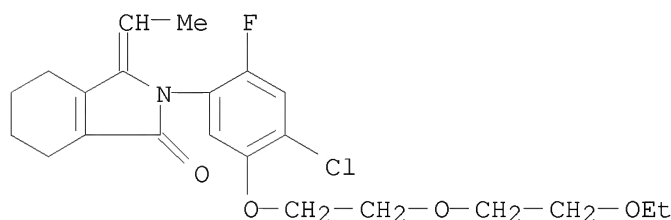
RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of)

RN 133715-52-3 CAPLUS

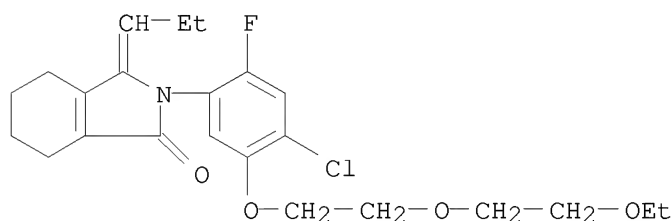
CN 1H-Isindol-1-one, 2-[4-chloro-3-(2-propen-1-ylthio)phenyl]-3-ethylidene-  
2,3,4,5,6,7-hexahydro- (CA INDEX NAME)



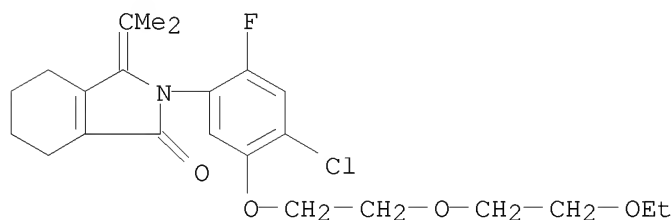
RN 133715-56-7 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-5-[2-(2-ethoxyethoxy)ethoxy]-2-fluorophenyl]-3-ethylidene-2,3,4,5,6,7-hexahydro- (CA INDEX NAME)



RN 133715-58-9 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-5-[2-(2-ethoxyethoxy)ethoxy]-2-fluorophenyl]-2,3,4,5,6,7-hexahydro-3-propylidene- (CA INDEX NAME)

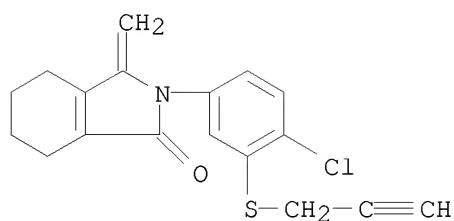


RN 133715-60-3 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-5-[2-(2-ethoxyethoxy)ethoxy]-2-fluorophenyl]-2,3,4,5,6,7-hexahydro-3-(1-methylethylidene)- (CA INDEX NAME)

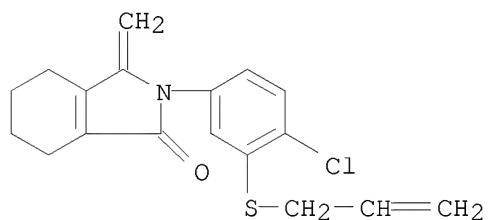


RN 133715-62-5 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-3-(2-propyn-1-ylthio)phenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)

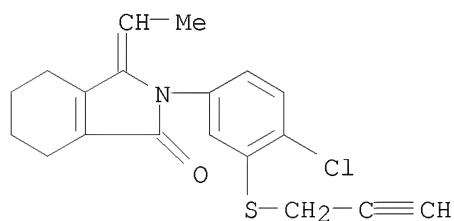




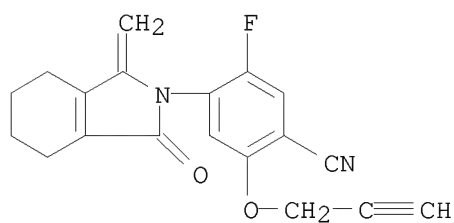
RN 133715-63-6 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-3-(2-propen-1-ylthio)phenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



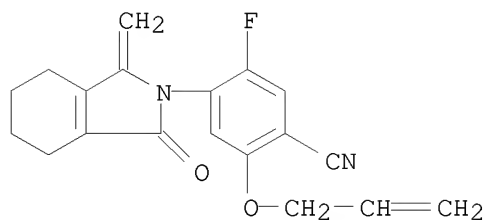
RN 133715-64-7 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-3-(2-propyn-1-ylthio)phenyl]-3-ethylidene-2,3,4,5,6,7-hexahydro- (CA INDEX NAME)



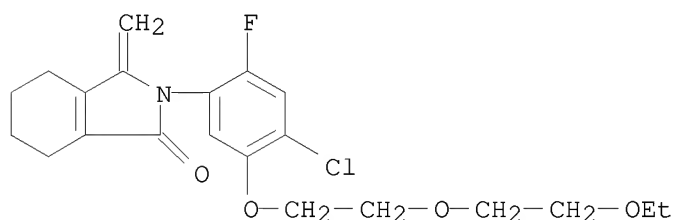
RN 133715-66-9 CAPLUS  
 CN Benzonitrile, 5-fluoro-4-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)-2-(2-propyn-1-yloxy)- (CA INDEX NAME)



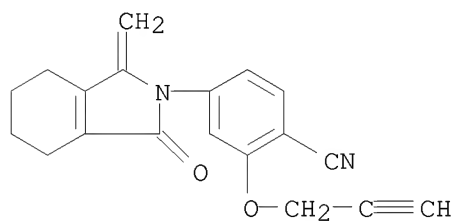
RN 133715-69-2 CAPLUS  
 CN Benzonitrile, 5-fluoro-4-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)-2-(2-propen-1-yloxy)- (CA INDEX NAME)



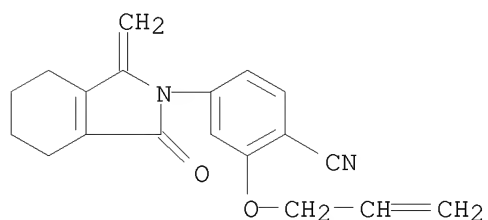
RN 133715-71-6 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-5-[2-(2-ethoxyethoxy)ethoxy]-2-fluorophenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



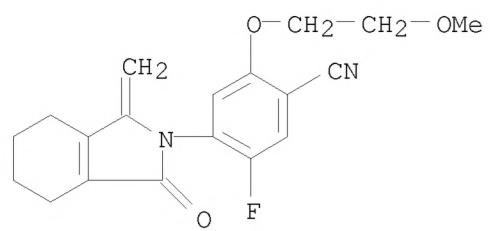
RN 133715-72-7 CAPLUS  
 CN Benzonitrile, 4-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)-2-(2-propyn-1-yloxy)- (CA INDEX NAME)



RN 133715-73-8 CAPLUS  
 CN Benzonitrile, 4-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)-2-(2-propen-1-yloxy)- (CA INDEX NAME)



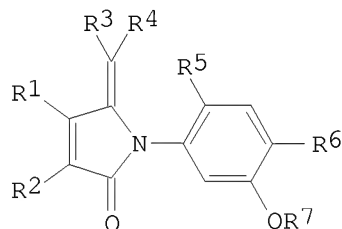
RN 133715-74-9 CAPLUS  
 CN Benzonitrile, 5-fluoro-4-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)-2-(2-methoxyethoxy)- (CA INDEX NAME)



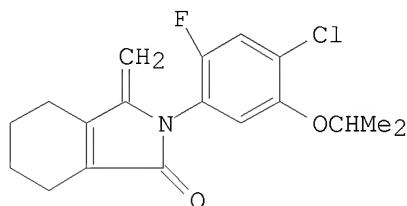
L4 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1989:533984 CAPLUS  
DOCUMENT NUMBER: 111:133984  
ORIGINAL REFERENCE NO.: 111:22426h, 22427a  
TITLE: Preparation and testing of 1-aryl-2-alkylidene-5-oxopyrrolines as herbicides and plant growth regulators  
INVENTOR(S): Pissiotas, Georg; Moser, Hans; Brunner, Hans Georg  
PATENT ASSIGNEE(S): Ciba-Geigy A.-G., Switz.  
SOURCE: Eur. Pat. Appl., 59 pp.  
CODEN: EPXXDW  
DOCUMENT TYPE: Patent  
LANGUAGE: German  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 305333	A2	19890301	EP 1988-810564	19880818
R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
AU 8821576	A	19890302	AU 1988-21576	19880826
BR 8804379	A	19890321	BR 1988-4379	19880826
HU 47540	A2	19890328	HU 1988-4468	19880826
HU 203081	B	19910528		
ZA 8806359	A	19890530	ZA 1988-6359	19880826
JP 01071855	A	19890316	JP 1988-213518	19880827
PRIORITY APPLN. INFO.:			CH 1987-3286	A 19870827
OTHER SOURCE(S):	MARPAT 111:133984			
GI				



I



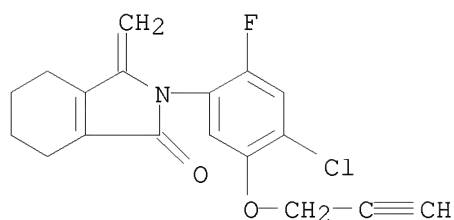
II

AB The title compds. [I; R1,R2 = H, C1-8 alkyl; R1R2 = (alkyl substituted) (CH2)4; R3,R4 = H, C1-4 alkyl, (substituted) Ph; R3R4 = (alkyl substituted) (CH2)n; n = 2-6; R5 = H, F; R6 = halo; R7 = R1, C3-8 alkenyl alkynyl, C3-7 cycloalkyl, substituted C1-4 alkyl], useful as herbicides, were prepared 2-(4-Chloro-2-fluoro-5-isopropoxyphenyl)-4,5,6,7-tetrahydroisindole-2H-1,3-dione in Et2O was added to MeMgI and the mixture was refluxed 1 h to give a 3° alc. which was stirred neat with KHSO4 for 30 min at 140° to give 2-aryl-3-methylene-tetrahydroisindole II. At 125 g/ha preemergent, II gave complete control of Echinochloa crus-galli and Monocharia vage in rice paddies, while leaving rice plants unaffected.

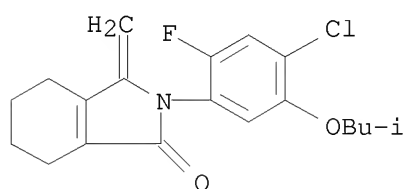
IT 115296-55-4P 122477-91-2P 122478-01-7P  
122478-02-8P 122478-05-1P  
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as herbicide and plant growth regulator)

RN 115296-55-4 CAPLUS

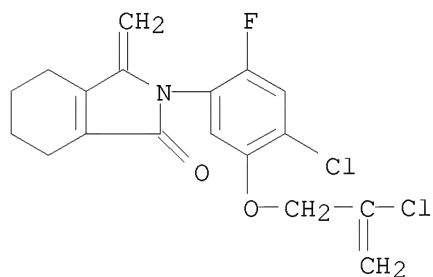
CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-propyn-1-yloxy)phenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



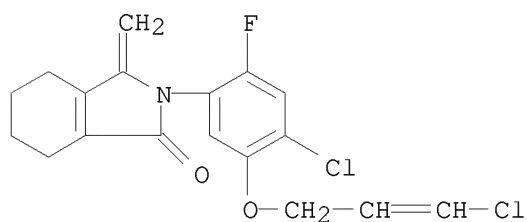
RN 122477-91-2 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-methylpropoxy)phenyl]-  
 2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



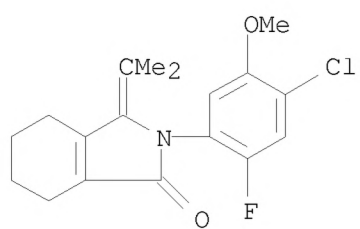
RN 122478-01-7 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-5-[(2-chloro-2-propen-1-yl)oxy]-2-  
 fluorophenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



RN 122478-02-8 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-5-[(3-chloro-2-propen-1-yl)oxy]-2-  
 fluorophenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



RN 122478-05-1 CAPLUS  
 CN 1H-Isoindol-1-one, 2-(4-chloro-2-fluoro-5-methoxyphenyl)-2,3,4,5,6,7-  
 hexahydro-3-(1-methylethylidene)- (CA INDEX NAME)

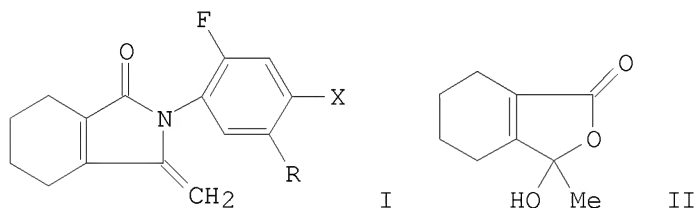


L4 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1988:437735 CAPLUS  
DOCUMENT NUMBER: 109:37735  
ORIGINAL REFERENCE NO.: 109:6383a,6386a  
TITLE: Preparation of tetrahydroisoindolone derivatives as herbicides  
INVENTOR(S): Saito, Kenji; Hagiwara, Kenji; Hara, Tamio; Ishikawa, Hisao; Inaba, Hideo; Sato, Junji  
PATENT ASSIGNEE(S): Nippon Soda Co., Ltd., Japan  
SOURCE: Jpn. Kokai Tokkyo Koho, 8 pp.  
CODEN: JKXXAF  
DOCUMENT TYPE: Patent  
LANGUAGE: Japanese  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63039859	A	19880220	JP 1986-182895	19860804
JP 07084442	B	19950913		
PRIORITY APPLN. INFO.:			JP 1986-182895	19860804
OTHER SOURCE(S):	MARPAT	109:37735		

GI

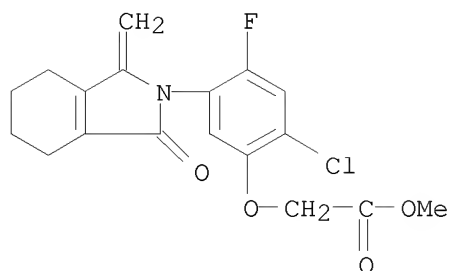


AB The title compds. I [X = halo; R = CO<sub>2</sub>R<sub>1</sub>, YR<sub>2</sub>; R<sub>1</sub> = H, (substituted) lower alkyl; Y = O, S; R<sub>2</sub> = H, (substituted) lower hydrocarbyl], useful as herbicides, were prepared from II. A mixture of II 4.5, p-MeC<sub>6</sub>H<sub>4</sub>SO<sub>3</sub>H 0.5, and 2-chloro-4-fluoro-5-aminophenol 4.33 g in 30 mL xylene was refluxed for 20 h to give 6.9 g I (X = Cl, R = OH) (III). At 5 g/are, III gave complete kill of *Cyperus difformis*.

IT 115296-45-2P 115296-46-3P 115296-52-1P  
115296-53-2P 115296-54-3P 115296-55-4P  
115296-56-5P 115296-60-1P 115296-61-2P  
115296-65-6P  
RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as herbicide)

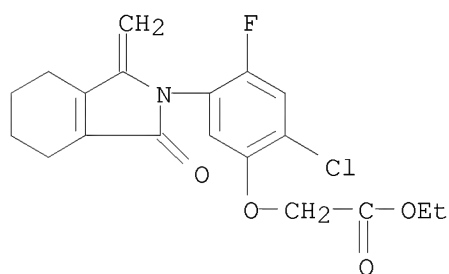
RN 115296-45-2 CAPLUS

CN Acetic acid, 2-[2-chloro-4-fluoro-5-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)phenoxy]-, methyl ester (CA INDEX NAME)



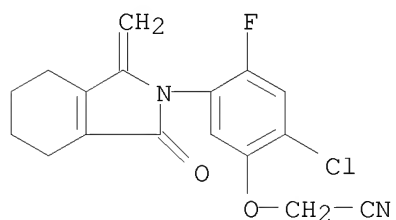
RN 115296-46-3 CAPLUS

CN Acetic acid, 2-[2-chloro-4-fluoro-5-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)phenoxy]-, ethyl ester (CA INDEX NAME)



RN 115296-52-1 CAPLUS

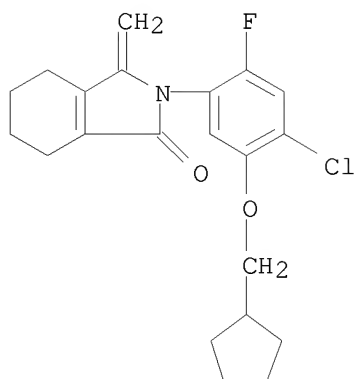
CN Acetonitrile, 2-[2-chloro-4-fluoro-5-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)phenoxy]- (CA INDEX NAME)



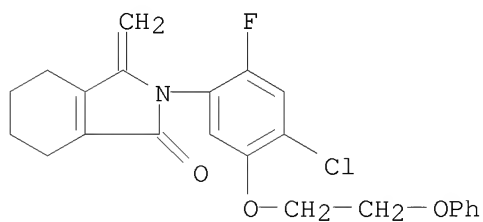
RN 115296-53-2 CAPLUS

CN 1H-Isoindol-1-one, 2-[4-chloro-5-(cyclopentylmethoxy)-2-fluorophenyl]-2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)

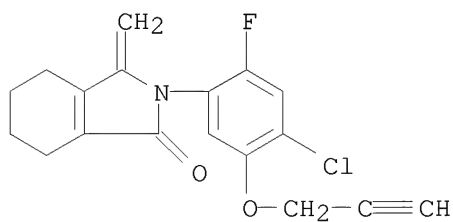




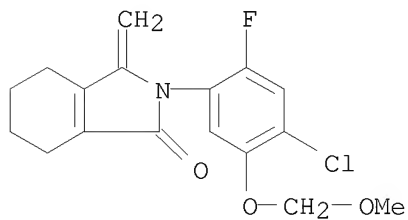
RN 115296-54-3 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-phenoxymethyl)phenyl]-  
 2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



RN 115296-55-4 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(2-propyn-1-yloxy)phenyl]-  
 2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)

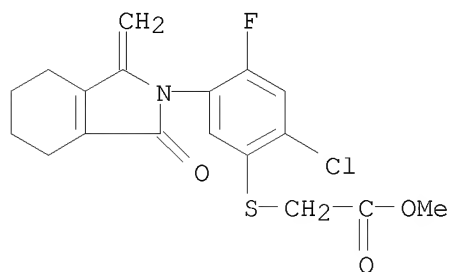


RN 115296-56-5 CAPLUS  
 CN 1H-Isoindol-1-one, 2-[4-chloro-2-fluoro-5-(methoxymethoxy)phenyl]-  
 2,3,4,5,6,7-hexahydro-3-methylene- (CA INDEX NAME)



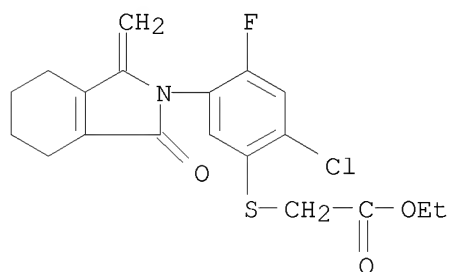
RN 115296-60-1 CAPLUS

CN Acetic acid, 2-[[2-chloro-4-fluoro-5-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)phenyl]thio]-, methyl ester (CA INDEX NAME)



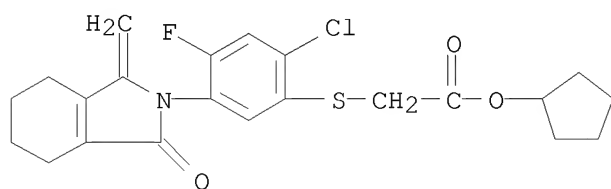
RN 115296-61-2 CAPLUS

CN Acetic acid, 2-[[2-chloro-4-fluoro-5-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)phenyl]thio]-, ethyl ester (CA INDEX NAME)



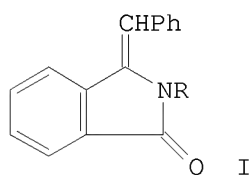
RN 115296-65-6 CAPLUS

CN Acetic acid, 2-[[2-chloro-4-fluoro-5-(1,3,4,5,6,7-hexahydro-1-methylene-3-oxo-2H-isoindol-2-yl)phenyl]thio]-, cyclopentyl ester (CA INDEX NAME)

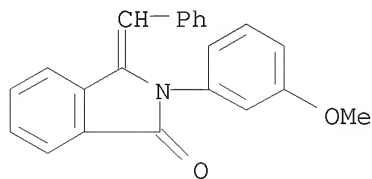


L4 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1980:181066 CAPLUS  
DOCUMENT NUMBER: 92:181066  
ORIGINAL REFERENCE NO.: 92:29333a,29336a  
TITLE: Studies on phthalimidines: Part I. Synthesis and fungitoxicity of some N-substituted phthalimidines  
AUTHOR(S): Mund, A. P.; Behera, R. K.; Nayak, A.; Behera, G. B.  
CORPORATE SOURCE: Postgrad. Dep. Chem., Sambalpur Univ., Sambalpur, 768017, India  
SOURCE: Indian Journal of Chemistry, Section B: Organic Chemistry Including Medicinal Chemistry (1979), 17B(4), 404-5  
CODEN: IJSBDB; ISSN: 0376-4699  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
GI



AB Twenty-one phthalimidines I [R = (un)substituted Ph or 2-benzothiazolyl] were prepared by reaction of benzaldehyde with the appropriate amine RNH<sub>2</sub>. I have marked fungitoxicity at 10 ppm.  
IT 73518-32-8P  
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)  
(preparation and fungicidal activity of)  
RN 73518-32-8 CAPLUS  
CN 1H-Isoindol-1-one, 2,3-dihydro-2-(3-methoxyphenyl)-3-(phenylmethylene)-  
(CA INDEX NAME)

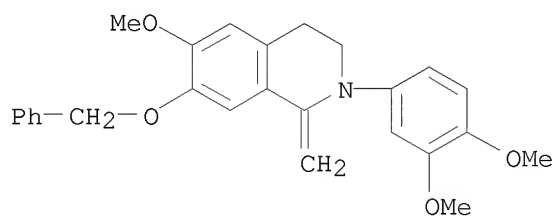


L4 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

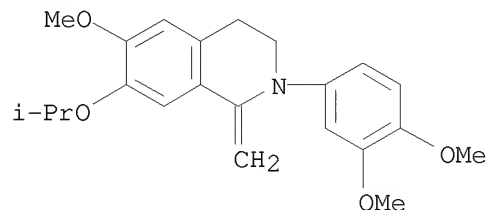
ACCESSION NUMBER: 1978:51054 CAPLUS  
DOCUMENT NUMBER: 88:51054  
ORIGINAL REFERENCE NO.: 88:8069a,8072a  
TITLE: Application of enamine photocyclization to a total  
synthesis of (±)-cryptaustoline  
AUTHOR(S): Ninomiya, Ichiya; Yasui, Junko; Kiguchi, Toshiko  
CORPORATE SOURCE: Kobe Women's Coll. Pharm., Higashinada, Japan  
SOURCE: Heterocycles (1977), 6(11), 1855-60  
CODEN: HTCYAM; ISSN: 0385-5414  
DOCUMENT TYPE: Journal  
LANGUAGE: English  
GI

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB (±)-Cryptaustoline (I) was prepared from the phenylacetanilide II, via  
photochem. cyclization of the anamine III to the dibenzopyrrocoline IV.  
IT 65341-63-1P  
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT  
(Reactant or reagent)  
(preparation and photochem. cyclization of, indoloisoquinoline derivs. from)  
RN 65341-63-1 CAPLUS  
CN Isoquinoline, 2-(3,4-dimethoxyphenyl)-1,2,3,4-tetrahydro-6-methoxy-1-  
methylene-7-(phenylmethoxy)- (CA INDEX NAME)



IT 65374-44-9P  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation and total cyclization of, indoloisoquinoline from)  
RN 65374-44-9 CAPLUS  
CN Isoquinoline, 2-(3,4-dimethoxyphenyl)-1,2,3,4-tetrahydro-6-methoxy-1-  
methylene-7-(1-methylethoxy)- (CA INDEX NAME)

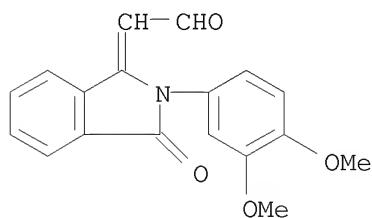


L4 ANSWER 10 OF 10 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1970:498642 CAPLUS  
DOCUMENT NUMBER: 73:98642  
ORIGINAL REFERENCE NO.: 73:16087a,16090a  
TITLE: 3-Formylmethylenephthalimidines  
INVENTOR(S): Seefelder, Matthias; Mueller, Hans Richard  
PATENT ASSIGNEE(S): Badische Anilin- & Soda-Fabrik AG  
SOURCE: Ger. Offen., 7 pp.  
CODEN: GWXXBX  
DOCUMENT TYPE: Patent  
LANGUAGE: German  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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DE 1905833	A	19700903	DE 1969-1905833	19690206
CH 542203	A	19731115	CH 1970-1291	19700129
GB 1288685	A	19720913	GB 1970-1288685	19700205
FR 2034016	A6	19701204	FR 1970-4213	19700206

PRIORITY APPLN. INFO.: DE 1969-1905833 A 19690206  
GI For diagram(s), see printed CA Issue.  
AB The title compds. (I) (R = CHO) were prepared from I (R = H) and R2R3NCHO (II). Thus, a mixture of I (R = H, R1 = p-O2NC6H4) in CHCl3, II (R2 = R3 = Me), and COCl2 was kept 1 hr at 0-5, 1 hr at 25, and 1 hr at 45° to give 90% I (R = CHO, R1 = p-O2NC6H4). Similarly prepared were I (R = CHO) (R1 given): p-ClC6H4; 2,4-(MeO)2C6H3; HO2CCH2, 2,4-Cl2C6H3, Et, iso-Pr.  
IT 29262-73-5P  
RL: SPN (Synthetic preparation); PREP (Preparation)  
(preparation of)  
RN 29262-73-5 CAPLUS  
CN  $\Delta 1, \alpha$ -Isoindolineacetaldehyde, 2-(2,4-dimethoxyphenyl)-3-oxo-  
(8CI) (CA INDEX NAME)



=> log y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

57.38

235.95

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

ENTRY

SESSION

CA SUBSCRIBER PRICE

-8.00

-8.00

STN INTERNATIONAL LOGOFF AT 10:34:20 ON 17 SEP 2008